AN ADC SP F- adenosine solution
AN Co Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, click here.

ACTIVE INGREDIENT

Active Ingredient: Adenosine 0.04%

INACTIVE INGREDIENT

SP F I) Inactive Ingredients: Water, Carbomer, Beta-Glucan, Caffeine, Avena Sativa (Oat) Kernel Extract, Hovenia Dulcis Fruit Extract, Centella Asiatica Extract/Diospyros Kaki Leaf Extract/Theobroma Cacao Extract/Chamomilla Recutita (Matricaria) Extract/Wine Extract, Glycine Soja (Soybean) Seed Extract, Origanum Vulgare Flower Extract, Octanediol (Carprylyl Glycol), Phenoxyethanol, Phenoxyethanol, Sophora Flavescens Root Extract, Scutellaria Baicalensis Root Extract, Acetyl Hexapeptide-8, Copper Tripeptide-1

SP F II) Inactive Ingredients: Sodium Silicate / Hydroxypropyl Chitosan, Water, Ipomoea Purpurea Extract/Paeonia Albiflora Flower Extract/Magnolia Liliflora Flower Extract/Lilium Candidum Flower Extract, Sodium Hydroxide, Camellia Sinensis Leaf Extract, Portulaca Oleracea Extract, Polyepsilon-Lysine

PURPOSE

Purpose: Skin Protectant

WARNINGS

Precautions on use: 1. Stop using the product if there are any of the following abnormal symptoms appearing after use and consult with your dermatologist as continued use can worsen the symptoms. A) If there are red spots, swelling, itchiness, or irritation B) If the above symptoms appear around the skin to which the product has been applied after being exposed to direct sunlight. 2. Precautions on storage and handling A) Make sure to close the cap after use. B) Keep out of the reach of children. C) Do not keep the product in a hot or cold place or a place getting direct sunlight. 3. Do not apply the product to any parts of the skin with wound, eczema, or dermatitis.

KEEP OUT OF REACH OF CHILDREN

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Suggested use and application order

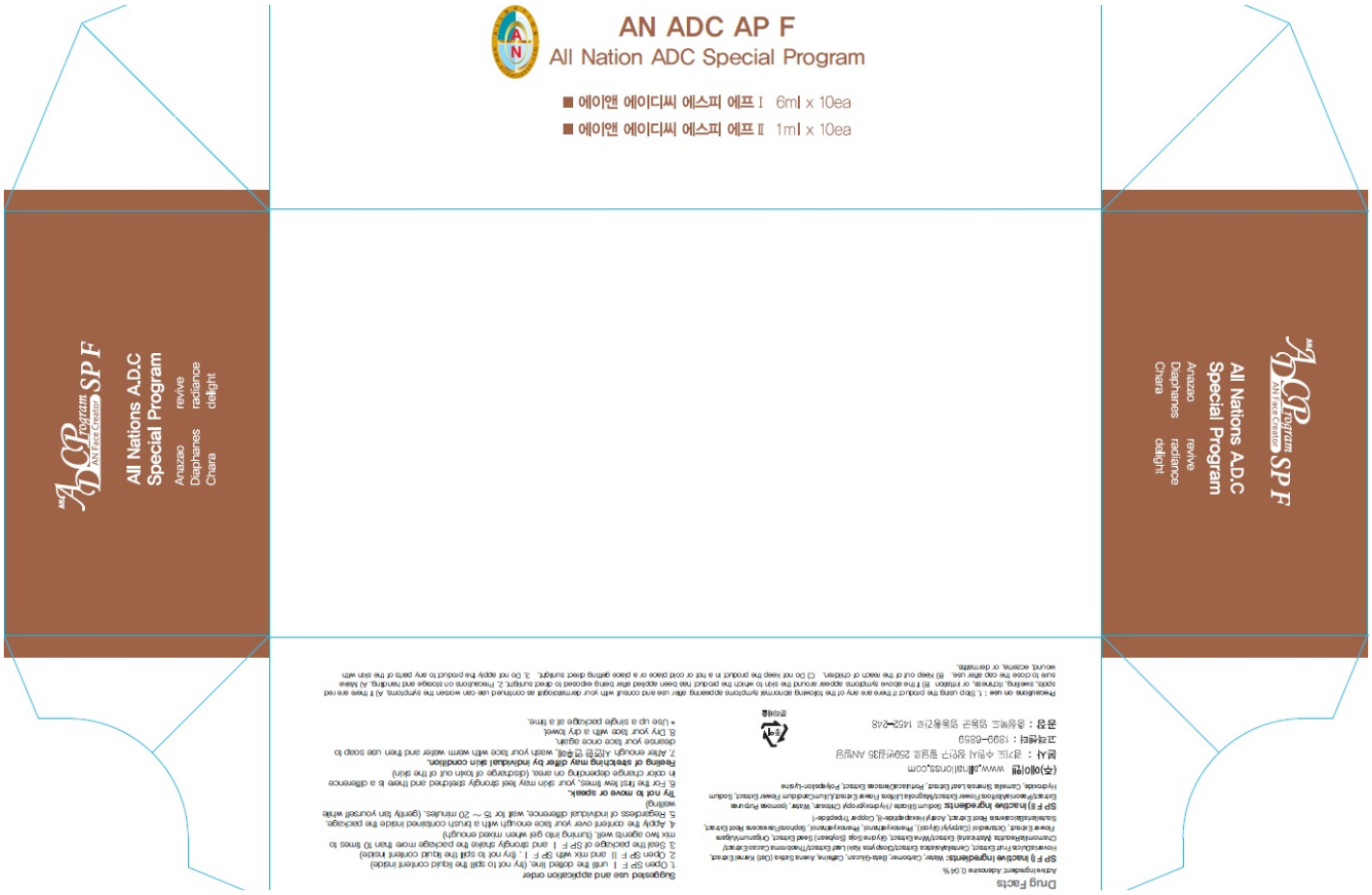
Suggested use and application order: 1. Open SP F □ until the dotted line. (try not to spill the liquid content inside) 2. Open SP F □ and mix with SP F □. (try not to spill the liquid content inside) 3. Seal the package of SP F □ and strongly shake the package more than 10 times to mix two agents well. (turning into gel when mixed enough) 4. Apply the content over your face enough with a brush contained inside the package. 5. Regardless of individual difference, wait for 15 ~ 20 minutes. (gently fan yourself while waiting) Try not to move or speak. 6. For the first few times, your skin may feel strongly stretched and there is a difference in color change depending on area. (discharge of toxin out of the skin) Feeling of stretching may differ by individual skin condition. 7. After enough application, wash your face with

warm water and then use soap to cleanse your face once again. 8. Dry your face with a dry towel.

DOSAGE & ADMINISTRATION

Dosage & administration: Use up a single package at a time.

PACKAGE LABEL.PRINCIPAL DISPLAY PANEL



AN ADC SP F

adenosine solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69153-070
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
Adenosine (UNII: K72T3FS567) (ADENOSINE - UNII:K72T3FS567)	Adenosine	2.8 mg in 7 mL

Inactive Ingredients	
Ingredient Name	**Strength**
Water (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69153-070-02	10 in 1 CARTON	05/01/2016	
1	NDC:69153-070-01	7 mL in 1 CONTAINER; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved drug other		05/01/2016	

Labeler - AN Co Ltd. (688448454)

Registrant - AN Co Ltd. (688448454)

Establishment

Name	Address	ID/FEI	Business Operations
AN Co Ltd.		688448454	manufacture(69153-070)

Revised: 6/2016 AN Co Ltd.